Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1. (original) A compound of formula

$$R_{1}$$

wherein

R₁ is halogen or halo(C₁₄)alkyl,

R₂ is hydrogen, halogen or halo(C₁₋₄)alkyl,

R₃ is halogen or halo(C₁₋₄)alkyl,

R₄ is hydrogen, (C₁₋₈)alkyl, hydroxy(C₁₋₆)alkyl or a group of formula

- -CO-R₅,
- -CO-(CH₂)_m-OR₆,
- -CO-CO-R₇,
- -CO-CO-OR₈,
- -CO-N(R₉R₁₀),
- -CO-(CH₂)_n-CO-R₁₁,
- -CO-(CHR₁₅)-O-(CH₂)_o-CO-R₁₁,
- $-CO-(CH_2)_p-O-(CH_2)_q-O-(CH_2)_r-R_{16}$
- -CO-O-(CH₂)_s-O-CO-R₁₇,
- $-CO-O-(CH_2)_t-N(R_{18}R_{19}),$
- -CO-O- $(CH_2)_u$ -NH-CO- $CH(NH_2)$ - R_{20} , or
- -CO-O-(CH₂)_w-NH-CO-R₁₇, wherein
 - R₅ is hydrogen, (C₁₋₈)alkyl, (C₃₋₈)cycloalkyl, amino, (C₁₋₄)alkylamino, di(C₁₋₄)alkylamino, aryl or heterocyclyl which is a 5 or 6-membered heterocyclic ring system having 1 to 4 heteroatoms selected from N, O or S,
 - R₆ is hydrogen, (C₁₋₄)alkyl, (C₃₋₈)cycloalkyl, aryl, (C₁₋₄)alkyl substituted by heterocyclyl which is a 5 or 6-membered heterocyclic ring system having 1 to 4 heteroatoms selected from N, O or S, amino(C₁₋₆)alkyl,
 - (C_{1-4}) alkylamino (C_{1-6}) alkyl, di (C_{1-4}) alkylamino (C_{1-6}) alkyl, hydroxy (C_{1-6}) alkyl,

hydroxy(C_{1^-4})alkylamino(C_{1^-6})alkyl or an amino acid residue, e.g. $-CH_2$ - $CH(NH_2)$ -COOH,

R₇ and R₈ independently of each other are (C₁₋₄)alkyl, (C₃₋₈)cycloalkyl, aryl or heterocyclyl which is a 5 or 6-membered heterocyclic ring system having 1 to 4 heteroatoms selected from N, O or S,

 R_9 and R_{10} independently of each other are hydrogen or (C_{1-4}) alkyl or one of R_9 and R_{10} is hydrogen and the other is (C_{3-8}) cycloalkyl, (C_{1-4}) alkyl, aryl or heterocyclyl,

 R_{11} is (C_{1-4}) alkyl, $-OR_{12}$, $-NR_{13}R_{14}$, an amino acid, an (C_{1-4}) alkylester thereof or a di (C_{1-4}) alkylester thereof,

R₁₂ is hydrogen or (C₁₋₄)alkyl,

 R_{13} and R_{14} independently of each other are hydrogen, (C_{1-4}) alkyl, amino (C_{1-6}) alkyl, (C_{1-4}) alkylamino (C_{1-6}) alkyl, di (C_{1-4}) alkylamino (C_{1-6}) alkyl,

R₁₅ is hydrogen or (C₁₋₄)alkyl,

R₁₆ is hydrogen, (C₁₋₄)alkyl, carboxyl or carboxylic ester,

 R_{17} is amino(C_{1-4})alkyl, (C_{1-4})alkylamino(C_{1-4})alkyl or di(C_{1-4})alkylamino(C_{1-4})alkyl,

R₁₈ is hydrogen or (C₁₋₄)alkyl,

 R_{19} is hydroxy(C_{1-4})alkyl,

 R_{20} is (C_{1-4}) alkyl or hydroxy (C_{1-4}) alkyl,

m is 0 to 4,

n is 2 to 8.

o is 0 to 4,

p is 0 to 4,

q is 1 to 8,

r is 0 to 4,

s is 1 to 4,

t is 1 to 4,

u is 1 to 6 and

w is 1 to 6.

Claim 2. (original) A compound of claim 1 wherein

- R₁ is chloro or trifluoromethyl,
- R₂ is hydrogen or trifluoromethyl,
- R₃ is chloro, fluoro or trifluoromethyl,
- R_4 is hydrogen, (C_{1-4}) alkyl, e.g. methyl, hydroxy (C_{1-4}) alkyl, e.g. hydroxyethyl, or a group of formula
 - -CO-R₅,
 - -CO-(CH₂)_m-OR₆,

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-CO-CO-R7.
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- -CO-CO-OR₈,
- -CO-N(R₉R₁₀),
- -CO-(CH₂)_n-CO-R₁₁,
- -CO-(CHR₁₅)-O-(CH₂)₀-CO-R₁₁,
- -CO-(CH₂)_p-O-(CH₂)_q-O-(CH₂)_r-R₁₆,
- -CO-O-(CH₂)_s-O-CO-R₁₇,
- -CO-O-(CH₂)_t-N(R₁₈R₁₉),
- -CO-O-(CH₂)_u-NH-CO-CH(NH₂)-R₂₀, or
- -CO-O-(CH₂)_w-NH-CO-R₁₇, wherein

 R_5 is hydrogen, (C_{1-4}) alkyl, (C_{3-6}) cycloalkyl, dimethylamino, phenyl or heterocyclyl which is a 6-membered heterocyclic ring system having one O as a heteroatom, e.g. tetrahydropyranyl,

R₆ is hydrogen, (C₁₋₄)alkyl, (C₁₋₂)alkyl substituted by heterocyclyl which is a 5 or 6-membered heterocyclic ring system having 1 or 2 heteroatoms selected from N or O, e.g. including unsubstituted pyrrolidine, morpholine and piperazine and piperazine substituted by e.g. (C₁₋₂)alkyl or (C₁₋₂)hydroxyalkyl; amino(C₁₋₄)alkyl, (C₁₋₂)alkylamino(C₁₋₄)alkyl, di(C₁₋₂)alkylamino(C₁₋₄)alkyl, hydroxy(C₁₋₃)alkyl, hydroxy(C₁₋₂)alkylamino(C₁₋₂)alkyl or an amino acid residue, e.g. –CH₂-CH(NH₂)-COOH,

R₇ and R₈ independently of each other are (C₁₋₂)alkyl or phenyl,

R₉ and R₁₀ independently of each other are hydrogen or (C₁₋₂)alkyl,

R₁₁ is (C_{1-2}) alkyl, $-OR_{12}$, $-NR_{13}R_{14}$, an amino acid, an (C_{1-2}) alkylester thereof or an di (C_{1-2}) alkylester thereof, preferably an amino acid selected from the group consisting of alanine, phenylalanine, glutamic acid and lysine, wherein the binding is effected via the α - amino group or in the case of e.g. lysine via the ϵ -amino group,

R₁₂ is hydrogen or (C₁₋₂)alkyl,

 R_{13} and R_{14} independently of each other are hydrogen, (C_{1-2}) alkyl, amino (C_{1-4}) alkyl, (C_{1-2}) alkylamino (C_{1-4}) alkyl, di (C_{1-2}) alkylamino (C_{1-4}) alkyl,

R₁₅ is hydrogen or (C₁₋₂)alkyl,

R₁₆ is hydrogen, (C₁₋₂)alkyl, carboxyl or carboxylic ester,

 R_{17} is amino(C_{1-2})alkyl,

 R_{18} is hydrogen or (C_{1-2}) alkyl,

 R_{19} is hydroxy(C_{1-2})alkyl,

 R_{20} is (C_{1-2}) alkyl or hydroxy (C_{1-2}) alkyl,

m is 0 or 1,

n is 2 to 4.

o is 0 or 1, p is 0 to 2, q is 2 to 5, r is 0 to 2, s is 2, t is 2, u is 1 to 3 and w is 1 to 3.

Claim 3. (currently amended) A compound according to claim 1-or 2 which is a compound of formula I wherein

R₁ is chloro,

R₂ is hydrogen,

R₃ is trifluoromethyl and

R₄ is hydrogen.

Claim 4. (currently amended) A compound according to claim 1-or 2 which is a compound of formula I wherein

R₁ is chloro,

R₂ is hydrogen,

R₃ is trifluoromethyl and

 R_4 is a group of formula $-CO-O-(CH_2)_2-N[(C_2H_5OH)(CH_3)]$.

Claim 5. (currently amended) A compound according to any one of claims 1-to 4 in the form of a salt.

Claim 6. (canceled).

Claim 7. (currently amended) A method of treatment of IgE-synthesis-mediated diseases, autoimmune diseases, gastrointestinal diseases and chronic rejection of transplants which method comprises administering a therapeutically effective amount of a compound of any one of claims 1-to-5 to a subject in need of such treatment.

Claim 8. (currently amended) A compound of any one of claims 1-to 5 for use as a pharmaceutical.

Claim 9. (currently amended) A pharmaceutical composition comprising a compound of any one of claims 1-to-5 in association with at least one pharmaceutical excipient.

Claim 10. (currently amended) Use of an amine, which is substituted by

- phenyl-substituted pyrimidin; and
- phenyl; and
- a third substituent, e.g. R_4 as defined in claim 1-to-5, in the preparation of a medicament for the treatment of IgE-synthesis-mediated diseases, autoimmune diseases, gastrointestinal diseases and chronic rejection of transplants.